PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

	cant's or agent's file reference [25622	FOR FURTHER AC	CTION	See Form PCT/IPEA/416
i	national application No.	International filing date (day/month/year)	Priority date (day/month/year) 19.05.2003
		C) or national classification and IF	PC	
ì	2N15/82, C07K14/01	Of or national olassinositives		
	······································			
Appl	icant	E TECNOLOGIE, L'ENERG	et al	
ENE	EA-ENTE PER LE NOOVE			
1.	This report is the internation Authority under Article 35 a	nal preliminary examination re and transmitted to the applican	port, established by t t according to Article	this International Preliminary Examining 36.
2.	This REPORT consists of a	a total of 6 sheets, including th	nis cover sheet.	
3.	This report is also accompa	anied by ANNEXES, comprisir	ng:	·
	a. 🛭 sent to the applican	t and to the International Bure	au) a total of 4 shee	ets, as follows:
	 sheets of the de and/or sheets or Administrative I 	ontaining rectifications authori	ngs which have been zed by this Authority	amended and are the basis of this report (see Rule 70.16 and Section 607 of the
		de parlier shoots but W	hich this Authority co	onsiders contain an amendment that goes
	Supplemental E	Box.		ndicated in item 4 of Box No. I and the
	Linking on	dertables related thereto in (ombliter readable to	nber of electronic carrier(s)) , containing a rm only, as indicated in the Supplemental
	Box Relating to Sec	quence Listing (see Section 80	2 of the Administrativ	ve instructions).
				·
	This report contains indica	tions relating to the following i	tems:	
4.				
	⊠ Box No. I Basis of □ □	the opinion		
			t inventi	ive eten and industrial applicability
			ard to noveity, invent	ive step and industrial applicability
	☐ Box No. IV Lack of u	unity of invention	a)	other inventive step or industrial
		ed statement under Afficie 35(ility; citations and explanation:	s supporting such sta	elty, inventive step or industrial Itement
		documents cited		
		defects in the international app		
	Box No. VIII Certain o	observations on the internation	nal application	
L				
Date	e of submission of the demand	·	Date of completion of	of this report
			47.00.0005	
18.	03.2005		17.08.2005	
Nan	ne and mailing address of the in	ternational	Authorized Officer	LAS PERS.
preliminary examining authority:			Logarition Margan E	
-	European Patent Office D-80298 Munich		Mundel, C	
	Tel. +49 89 2399 - 0 Fax: +49 89 2399 - 4	Tx: 523656 epmu d 465	Telephone No. +49	89 2399-7314
_	- 1 U.A. T-TO GO EGGO T-			

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

AP20 Recid Printernational application 2005 PCT/IT2004/000287

_	ox No. I Basis of the report				
1.	Ith regard to the language , this report is based on the international application in the language in which it was ed, unless otherwise indicated under this item.				
	This report is based on translations from the original language into the following language English, which is the language of a translation furnished for the purposes of:				
	 ☑ international search (under Rules 12.3 and 23.1(b)) ☐ publication of the international application (under Rule 12.4) ☐ international preliminary examination (under Rules 55.2 and/or 55.3) 				
2. With regard to the elements* of the international application, this report is based on (replacement she have been furnished to the receiving Office in response to an invitation under Article 14 are referred to report as "originally filed" and are not annexed to this report):					
	escription, Pages				
	as originally filed				
	Sequence listings part of the description, Pages				
	as originally filed				
	claims, Numbers				
	received on 30.03.2005 with letter of 18.03.2005				
	Prawings, Sheets				
	/16-16/16 as originally filed				
	a sequence listing and/or any related table(s) - see Supplemental Box Relating to Sequence Listing				
3.	The amendments have resulted in the cancellation of: ☐ the description, pages ☐ the claims, Nos. ☐ the drawings, sheets/figs ☐ the sequence listing (specify): ☐ any table(s) related to sequence listing (specify):				
4.	This report has been established as if (some of) the amendments annexed to this report and listed below had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)). the description, pages the claims, Nos. the drawings, sheets/figs the sequence listing (specify): any table(s) related to sequence listing (specify):				

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No. PCT/IT2004/000287

Box No. II Priority					
1.	\boxtimes	This report has been established as if no priority had been claimed due to the failure to furnish within the prescribed time limit the requested: ☑ copy of the earlier application whose priority has been claimed (Rule 66.7(a)). ☐ translation of the earlier application whose priority has been claimed (Rule 66.7(b)).			
2.		This report has been established as if no priority had been claimed due to the fact that the priority claim has been found invalid (Rule 64.1). Thus for the purposes of this report, the international filing date indicated above is considered to be the relevant date.			
3.	Add	litional observations, if necessary:			

Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

1-25 Yes: Claims Novelty (N) Claims No: Yes: Claims 1-25 Inventive step (IS) Claims No: 1-25 Claims Industrial applicability (IA) Yes: No: Claims

2. Citations and explanations (Rule 70.7):

see separate sheet

Box No. VIII Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate sheet

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No. PCT/IT2004/000287

_	Sup	ple	emental Box relating to Sequence Listing				
Co	Continuation of Box I, item 2:						
1.	With nec	n re	egard to any nucleotide and/or amino acid sequence disclosed in the international application and sary to the claimed invention, this report has been established on the basis of:				
	a. type of material:						
		⅓	a sequence listing				
]	table(s) related to the sequence listing				
	b. fo	orm	nat of material:				
	[X)	in written format				
		\boxtimes	in computer readable form				
	c. ti	me	of filing/furnishing:				
	0	\boxtimes	contained in the international application as filed				
	[filed together with the international application in computer readable form				
	[furnished subsequently to this Authority for the purposes of search and/or examination				
	[received by this Authority as an amendment on				
2.		th ac	addition, in the case that more than one version or copy of a sequence listing and/or table(s) relating ereto has been filed or furnished, the required statements that the information in the subsequent or diditional copies is identical to that in the application as filed or does not go beyond the application as filed, appropriate, were furnished.				
3.	Add	ditic	onal observations, if necessary:				

International application

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (SEPARATE SHEET)

PCT/IT2004/000287

Re Item V

Reasoned statement with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

- 1. The present application refers to a mutated V1/AR1/AV1 or C1/AL1/AC1 gene sequence of a tomato infecting geminivirus wherein the mutations consist of point mutations distributed along the sequence in such a way that the continuous homology between the mutated sequence and the corresponding viral gene sequence is below or equal to 8 nucleotides, said mutated sequence encoding for a capsid protein or for a Rep protein, to synthetic constructs comprising such a mutated gene sequence, to expression vectors comprising such constructs, to transgenic plants and seeds comprising such mutated gene sequence and to methods for the preparation of transgenic plants, plant tissues or cells thereof having long lasting resistance against geminiviruses.
- 2. New claims 1-25 filed with the letter of 17.03.2005 seem to comply with the requirements of Article 19(2) and 34(2)(b) PCT.
 - The remarks filed by the Applicant in the letter of 17.03.2005 have been taken into account for drafting the International Preliminary Examination Report (IPER).
- 3. Novelty and inventive step; Article 33(2) and 33(3) PCT.

In transgenic plants resistant to geminiviruses due to the expression of a viral protein, loss of resistance is observed after a certain time. None of the documents cited in the International Search Report (ISR) discloses or even suggests that the loss of resistance observed in the transgenic plants could be due to geminivirus-mediated silencing of the transgene. Therefore, the skilled person would not have been motivated to modify the sequence of the transgenes according to the present application.

Thus, claims 1-25 are to be considered as novel (Article 33(2) PCT) and inventive (Article 33(3) PCT).

Re Item VIII

Certain observations on the international application

- 1. Claim 1 refer to a mutated V1/AR1/AV1 or C1/AL1/AC1 gene sequence of a tomato infecting geminivirus encoding for a capsid protein or for a Rep protein. It is not clear if this protein is the native geminivirus protein or a modified protein and if the mutated gene encodes the same protein as the non-mutated gene or not. This renders the scope of claim 1 unclear.
- 2. In claim 3, there is no limit (minimal or maximal) to the size of the truncation what renders the scope of the claim unclear.
- 3. The wording of claim 5 is confuse. It is not clear if claim 5 is restricted to the sequences disclosed in SEQ ID NO: 3 and SEQ ID NO: 5 or if these sequences are only given as examples.

This remark also applies mutatis mutandis to claim 7.

- 4. Claim 12 b) refers to the mutagenesis of the viral sequence so as to make it an ineffective target of the post-transcriptional gene silencing induced by the infecting geminivirus. The attention of the Applicant is drawn that only two types of mutagenesis have been disclosed in the present application: (1) point mutations distributed along the sequence in such a way that the continuous homology between the mutated sequence and the corresponding viral gene sequence is below or equal to 8 nucleotides and (2) deletions of the 5' or 3' regions of the viral gene sequence of step a) until the identification of the region of said gene sequence that is an ineffective target of post-transcriptional gene silencing. Only these two types of mutations could be considered as fully supported by the description of the present application (Article 5 PCT when read in combination with Article 6 PCT).
- 5. In claims 15-17, it is not clear what is meant by "isolate thereof".

10

15

20

25

30

35

CLAIMS

- 1. Mutated V1/AR1/AV1 or C1/AL1/AC1 gene sequence of a tomato infecting geminivirus wherein the mutations consist of point mutations distributed along the sequence in such a way that the continuous homology between the mutated sequence and the corresponding viral gene sequence is below or equal to 8 nucleotides, preferably below or equal to 5 nucleotides, said mutated sequence encoding for a capsid protein or for a Rep protein, respectively.
- 2. Mutated V1/AR1/AV1 gene sequence according to claim 1, encoding for a capsid protein having sequence SEQ ID No 7.
- 3. Mutated C1/AL1/AC1 gene sequence according to claim 1, wherein the mutation further comprises a truncation occurring at 3' terminal so that the mutated sequence encodes for a truncated Rep protein.
- 4. Mutated C1/AL1/AC1 gene sequences according to claim 3, wherein the truncated Rep proteins consist of 130 aminoacids (Rep 130) to 210 aminoacids (Rep 210).
- 5. Mutated C1/AL1/AC1 gene sequence according to any of the claims 3 and 4 encoding for Rep 210 SEQ ID No 3 or SEQ ID No 5.
- 6. Mutated C1/AL1/AC1 gene sequence encoding for Rep 130 SEQ ID No 9.
- 7. Mutated gene sequence according to any of the claims 1-6 wherein the tomato infecting geminivirus is TYLCSV.
- 8. Synthetic construct comprising an heterologous polynucleotide sequence containing in the 5'-3' direction:
- a) polynucleotide sequence acting as promoter in said plant or tissue or transformed cells;
- b) a non translated polynucleotide sequence positioned 5' of the encoding region;
 - c) a mutated gene sequence according to any of the claims 1 to 7;
- d) a sequence acting as transcription terminator, positioned 3' with respect to the mutated gene sequence.
- Expression vector comprising the construct as defined according to claim 8.

10

15

20

25

30

35

38

- 10. Transgenic plant, tissue or plant cells thereof, comprising in their genome a mutated gene sequence according to any of the claims 1 to 7.
- 11. Seed comprising in its genome a mutated gene sequence according to any of the claims 1 to 7.
- 12. Method for the preparation of transgenic plants, plant tissue or cells thereof having long lasting resistance against geminiviruses, including the following steps:
- a) identification or selection of a viral gene sequence encoding an aminoacid sequence able to confer resistance against geminiviruses;
- b) mutagenesis of the viral gene sequence so as to make it an ineffective target of the post-transcriptional gene silencing induced by the infecting geminivirus;
- c) insertion of the geminivirus gene sequence mutated in the step b) in the plant, plant tissue or cell thereof, using a construct comprising an heterologous polynucleotide sequence containing in the 5'-3' direction:
- i) a polynucleotide sequence acting as promoter in said plant or tissue or transformed cells;
- ii) a non translated polynucleotide sequence positioned 5' of the encoding region;
- iii) a polynucleotide sequence encoding a geminivirus-derived aminoacid sequence, properly mutagenised to be an ineffective target of post-transcriptional gene silencing induced by the infecting geminivirus;
- iv) a sequence acting as transcription terminator positioned 3' with respect to said polynucleotide sequence
- 13. Method according to claim 12 wherein the mutations consist of point mutations distributed along the sequence in such a way that the continuous homology between the mutated sequence and the corresponding viral gene sequence is below or equal to 8 nucleotides, preferably below or equal to 5 nucleotides.
- Method according to claim 12, wherein the mutagenesis in step b) consists of deletions of the 5' or 3' regions of the viral gene sequence of step a) until the identification of the minimum region of said gene sequence that is an ineffective target of the post-transcriptional gene silencing induced by the infecting geminivirus compare to the original viral

10

15

20

25

30

35

39

sequence and that said truncated protein maintains the ability to confer resistance against geminiviruses.

- 15. Method according to any of the claims 12-14 wherein the geminiviruses are selected from the group consisting of species of Mastrevirus, Curtovirus, Begomovirus and Topocuvirus and isolates thereof.
- 16. Method according to claim 15, wherein Begomoviruses species are selected from the group consisting of TYLCCNV, TYLCGV, TYLCMalV, TYLCSV, TYLCTHV, TYLCV, ACMV, BGMV, CalCuV, ToCMoV, TGMV, ToGMoV, ToMHV, ToMoTV, ToMoV, ToRMV, ToSLCV, ToSRV, Cotton leaf curl (CLCrV, CLCuAV, ClCuGV, CLCuKV, CLCuMV, CLCuRV), East African cassava mosaic (EACMCV, EACMMV, EACMV, EACMZV), Potato yellow mosaic (PYMPV, PYMTV, PYMV), Squash leaf curl (SLCCNV, SLCV, SLCYV), Sweet potato leaf curl (SPLCGV, SPLCV), Tobacco leaf curl (TbLCJV, TbLCKoV, TbLCYNV, TbLCZV), Tomato leaf curl (ToLCBV, ToLCBDV, ToLCGV, ToLCKV, ToLCLV, ToLCMV, ToLCNDV, ToLCSLV, ToLCTWV, ToLCV, ToLCV) and isolates thereof.
- 17. Method according to claim 15, wherein the species belonging to the genus Mastrevirus, Curtovirus, Topocuvirus are selected from the group consisting of WDV, MSV, SSV, BYDV, TYDV, BCTV and isolates thereof.
- 18. Method according to any of the claims 12-17, wherein the gene sequence is selected from the group consisting of C1/AL1/AC1, C2/AL2/AC2, C3/AL3/AC3, C4/AL4/AC4, V1/AR1/AV1, V2/AR2/AV2, BC1/BL1 and BV1/BR1, belonging to the geminiviruses.
- 19. Method according to claim 18, wherein the C1/AL1/AC1 gene sequence belongs to TYLCSV.
- 20. Method according to claims 12 and 19, wherein the aminoacid sequence is a truncated protein with respect to the viral wild-type protein.
- 21. Method according to claims 12-14 and 20 wherein the viral gene sequences made ineffective targets of the virus-induced post-transcriptional gene silencing are the SEQ ID No 8, SEQ ID No 2 and SEQ ID No 4.
- 22. Method according to claim 21, wherein the truncated proteins are Rep-130 (SEQ ID No 9) or Rep-210 (SEQ ID No 3 and 5).
- 23. Method according to claim 18, wherein the V1/AR1/AV1 gene sequence belongs to TYLCSV

40

- 24. Method according to claims 13 and 23 wherein the viral gene sequence made an ineffective target of the virus-induced post-transcriptional gene silencing is the SEQ ID No 6 encoding for the capsid protein SEQ ID No 7.
- 25. Method according to anyone of the claims 12-24, wherein the plants, tissues or cells thereof belong to the group consisting of tomato, pepper, tobacco, squash, manioc, sweet potato, cotton, melon, potato, soybean, corn, wheat, sugar cane, bean, beet.